

K162199 Cascade IOMAX Intraoperative MonitorMar 2, 2017
209 days to decisionK162199 · Product code: **GWF** · Neurology
Source: <https://www.510kdatabase.net/k162199/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Stimulator, Electrical, Evoked Response (GWF) |
| Date received | Aug 5, 2016 |
| Decision date | Mar 2, 2017 |
| Days to decision | 209 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Cadwell Industries, Inc. |
| Location | Kennewick, WA, US |
| Contact | MELISSA KIRKUP |
| 510(k) history | 12 submissions · 12 cleared · 2014-2025 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k162199/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 24, 2026